Search History

DATE: Sunday, March 09, 2003 Printable Copy Create Case

Set Name Side by side Query		Hit Count	Set Name result set
DB=US	SPT,PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=OR		
<u>L9</u>	adenovir\$ near5 (CMV or cytomegalovir\$ or RSV or rous or MMTV or mouse near mammary) near10 muscle near5 (express\$ or prolong\$ or extend\$)	2	<u>L9</u>
<u>L8</u>	adenovir\$ near5 (CMV or cytomegalovir\$ or RSV or rous or MMTV or mouse near mammary) and muscle near5 (express\$ or prolong\$ or extend\$)	516	<u>L8</u>
<u>L7</u>	adenovir\$ near5 (CMV or cytomegalovir\$ or RSV or rous or MMTV or mouse near mammary) and muscle	1867	<u>L7</u>
<u>L6</u>	L4 and muscle near5 express\$ near10 (extend\$ or prolong\$ or durat\$)	12	<u>L6</u>
<u>L5</u>	L4 and muscle near5 express\$	925	<u>L5</u>
<u>L4</u>	L1 and (RSV or rous near sarcoma or CMV or cytomegalovirus or MMTV) near5 promoter\$	925	<u>L4</u>
<u>L3</u>	11 and RSV near5 promoter\$	313	<u>L3</u>
<u>L2</u>	adenovir\$ near5 vector\$ and muscle near5 express\$ near5 (prolong\$ or extend\$ or persist\$)	38	<u>L2</u>
<u>L1</u>	adenovir\$ near5 vector\$ and muscle near5 express\$	1320	<u>L1</u>

END OF SEARCH HISTORY

3/9/03 4:23 PM

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Search Results - Record(s) 1 through 12 of 12 returned.

1. 20030039657. 14 Apr 00. 27 Feb 03. Inducible vaccines. Johnston, Stephen Albert, et al. 424/184.1; 424/93.21 A61K048/00 A61K039/00.		
2. 20020037867. 26 Feb 99. 28 Mar 02. METHOD FOR RECOMBINANT ADENO-ASSOCIATED VIRUS-DIRECTED GENE THERAPY. WILSON, JAMES M., et al. 514/44; 435/320.1 435/455 435/456 435/457 435/69.1 A61K048/00 C12N015/861.		
3. 20010023349. 09 Apr 01. 20 Sep 01. Hypodermic needle with weeping tip and method of use. VanTassel, Robert A., et al. 606/53; A61B017/56.		
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11. 5780448 . 04 Nov 96; 14 Jul 98. DNA-based vaccination of fish. Davis; Heather L 514/44; 424/199.1 424/201.1 424/202.1 424/204.1 424/227.1 424/817 424/93.1 435/320.1 435/69.3 435/69.4 435/69.5 536/23.1 536/23.4 536/23.72. C12N015/00 A61K039/12 A61K039/29 A01N043/04.		
12. <u>5679647</u> . 03 Nov 94; 21 Oct 97. Methods and devices for immunizing a host against tumor-associated antigens through administration of naked polynucleotides which encode tumor-associated antigenic peptides. Carson; Dennis A., et al. 514/44; 424/184.1 536/23.1. A61K048/00 C12N015/12 C12N015/52.		
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         189884 ADENOVIR?
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          34858 ADENOVIR?(N) VECTOR?
        2782271 MUSCLE
         6890336 EXPRESS?
         151420 MUSCLE (5N) EXPRESS?
          25377 RSV
          30099 ROUS
             27 ADENOVIR? (N) VECTOR? AND MUSCLE (5N) EXPRESS? AND (RSV
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     S2 17 RD S1 (unique items)
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Processed 10 of 35 files ...
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     S5
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? rd s5
>>>Record 266:217889 ignored; incomplete bibliographic data, not retained -
in RD set
>>>Record 266:205079 ignored; incomplete bibliographic data, not retained -
in RD set
...completed examining records
             9 RD S5 (unique items)
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? d s6/9/1-9(Item 1 from file: 266) Display 6/9/1 DIALOG(R) File 266: FEDRIP Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv.

00341765

IDENTIFYING NO.: 5P01HL56091-05 0002 AGENCY CODE: CRISP ROLE OF CYTOMEGALOVIRUS IN AGA OF THE CORONARY ARTERY

PRINCIPAL INVESTIGATOR: HAYWARD, GARY

ADDRESS: JOHNS HOPKINS UNIV SCH OF MED 720 RUTLAND AVE BALTIMORE, MD 21205-2196

PERFORMING ORG.: JOHNS HOPKINS UNIVERSITY, BALTIMORE, MARYLAND SPONSORING ORG.: NATIONAL HEART, LUNG, AND BLOOD INSTITUTE FY: 2001 TYPE OF AWARD: Noncompeting Continuation (Type 5)

SUMMARY: Epidemiological evidence suggests that cytomegalovirus (HCMV) infection is a major contributing risk factor to the development of accelerated graft arteriosclerosis (AGA) in heart transplant recipients. We have obtained evidence for non-lytic chronic or latent infection by in situ hybridization in up to 70% of coronary artery sections in $\overline{\text{CMV}}$ positive recipients exhibiting AGA. HCMV IE1 and IE2 RNA, but not the 2.7kb IR4 delayed-early RNA were found to be expressed in large numbers of

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Display 6/9/1 (Item 1 from file: 266) DIALOG(R) File 266: FEDRIP Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv. infiltrating mononuclear lymphocytes or monocytes (MNC) and in many endothelial cells (EC) as well as in some clustered smooth muscle cells (SMC) in the intima. Patterns of HCMV IEl and IE2 protein expression correlated with the RNA results but differed in each cell type with SMC predominantly expressing the IE1 protein; EC the IE2 protein and MNC synthesizing neither protein. No late proteins were detectable except in occasional mature macrophages in the myocardium with inclusion bodies. We hypothesize that the expression of a limited subset of viral transcriptional regulatory genes in these non-permissively infected cells alter cellular gene expression and functional activity or the proliferation state of one or more of these cell types in ways that may trigger or exacerbate the changes involved in coronary arteriosclerosis. The specific aims of this project are thus to 1) Definitively characterize the in situ status of viral regulatory gene expression in different cell types in vessel walls with and without disease. 2) Examine the effects of infection and IE1or IE2 expression from adenovirus vectors on cellular levels of p53, TGF-beta and PDGF, in quiescent

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Display 6/9/1 (Item 1 from file: 266) DIALOG(R) File 266: FEDRIP Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv. and non-permissive BC and SMC culture systems with a view towards understanding the mechanisms involved in relevant cellular changes. We will also focus on the mechanisms of CMV.induced over-expression of p53 and determine whether the absence of p53 genes affects AGA in the presence or absence of CMV in a transgenic knock-out mouse model. 3) Examine otherwise inaccessible early events (especially in proliferating SMC in the intima) in a latently infected donor rat model that produces AGA within 20 days after cardiac transplantation and utilize genetically marked rat CMV to follow reactivation events. Parts of the work will be carried out in collaboration with other groups in the project. Our anticipation is that detailed knowledge of the role of CMV and its close relatives HHV6 and HHV.7 in AGA will lead to a better understanding of how to manage and monitor the disease, as well as to the development of potentially useful therapeutic or preventative strategies.

DESCRIPTORS: laboratory mouse; laboratory rat; transgenic animal; arteriosclerosis; vascular smooth muscle; vascular endothelium; cell type; pathologic process; tumor suppressor gene; gene expression; virus

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DIALOG(R) File 266: FEDRIP

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DIALOG(R)File 266:FEDRIP

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00302677

IDENTIFYING NO.: 5P01AR45925-03 0001 AGENCY CODE: CRISP ADENO-ASSOCIATED VIRUS (AAV) **VECTORS** TO IMPROVE MATURE **MUSCLE** FUNCTION

PRINCIPAL INVESTIGATOR: XIAO, XIAO

ADDRESS: UNIVERSITY OF PITTSBURGH PHARMACEUTICAL SCIENCES PITTSBURGH, PA 15213

PERFORMING ORG.: UNIVERSITY OF PITTSBURGH AT PITTSBURGH, PITTSBURGH, PENNSYLVANIA

SPONSORING ORG.: NAT INST OF ARTHRITIS AND MUSCULOSKELETAL AND SKIN DISEASES

FY: 2001 TYPE OF AWARD: Noncompeting Continuation (Type 5)

SUMMARY: Muscular dystrophies are a relatively common group of inherited degenerative **muscle** disease. Most types are caused by mutations in genes coding for membrance associated proteins in **muscle**. Duchenne muscular dystrophy (DMD) and limb-girdle muscular dystrophy (LGMD) often

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Display 6/9/2 (Item 2 from file: 266) DIALOG(R)File 266:FEDRIP

Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv. manifest themselves in young ages and lead to early morbidity with no currently available effective treatment. These diseases are recessive, loss-of- function of the corresponding gene product, which makes them suitable for gene replacement therapy. Recombinant adeno-associate virus (rAAV) is one promising gene replacement vector based on defective human parvoviruses. The rAAV system has attracted attention due to its non-pathogenicity, genomic integration, transduction of quiescent cells, and apparent lack of cellular immune reactions. In contrast to other viral vectors, rAAV is capable of efficiently bypassing the myofiber basal lamina and transducing mature muscle cells.

We have demonstrated that rAAV vectors harboring a foreign gene can achieve highly efficient and sustained gene expression in mature muscle of immunocompetent animals for more than 1.5 years without detectable toxicity. Recently, significant improvement in vector production methodology has made it possible to generate high titer and high quality rAAV vectors completely free of helper adenovirus contamination. However, no experiments using rAAV vectors to

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Display 6/9/2 (Item 2 from file: 266) DIALOG(R)File 266:FEDRIP

Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv. restore the functional deficits in muscle tissue itself have been reported to date. Here, we propose to take advantage of rAAV vector system, to test two therapeutic genes (delta-sarcoglycan and a highly truncated dystrophin), under the control of two different promoter systems (viral/CMV or muscle - specific/MCK), in two relevant animal models of muscular dystrophies (Biol4.6 hamster for LGMD and mdx mouse for DMD). Two distinct vector delivery methods, local intramuscular infection versus systemic delivery will be utilized. We have the following hypotheses to be tested. 1): muscle deficient in delta-sarcoglycan can be functionally rescued by genetic complementation using intramuscular AAV vector injection in the LGMD hamster model. systemic delivery of the delta-sarcoglycan gene can be mediated by rAAV vectors through intra-artery or intra-ventricle injection. 3) a dystrophin mini-gene lacking the central rod domain will improve the when delivered into function of dystrophin-deficient muscle dystrophic mdx mice by AAV vectors. DESCRIPTORS: hamster; laboratory mouse; intraarterial administration; -more-Display 6/9/2 (Item 2 from file: 266) DIALOG(R) File 266: FEDRIP Comp & dist by NTIS, Intl Copyright All Rights Res. All rts. reserv. qene therapy; gene complementation; muscular dystrophy; disease /disorder model; muscle protein; nonhuman therapy evaluation; adeno associated virus group; technology /technique development; dystrophin; transfection /expression vector; intramuscular injection - end of record -Display 6/9/3 (Item 1 from file: 434) DIALOG(R) File 434: SciSearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: G9232 Number of References: 31 08037271 Title: CDNA CLONING WITH A RETROVIRUS EXPRESSION VECTOR - GENERATION OF A PP60C-SRC CDNA CLONE Author(s): KAPLAN PL; SIMON S; CARTWRIGHT CA; ECKHART W Corporate Source: SALK INST BIOL STUDIES/SAN DIEGO//CA/92138 Journal: JOURNAL OF VIROLOGY, 1987, V61, N5, P1731-1734 Language: ENGLISH Document Type: NOTE Geographic Location: USA Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences Journal Subject Category: VIROLOGY Research Fronts: 86-1382 002 (HEMAGGLUTININ NEURAMINIDASE GENE OF NEWCASTLE-DISEASE VIRUS; NUCLEOTIDE-SEQUENCE ANALYSIS; TRANSCRIPTION OF GENES; CDNA CLONE) (TRANSCRIPTIONAL ENHANCER; VIRAL REGULATORY ELEMENTS; 86-1674 002 ADENOVIRUS E1A GENE; REGULATION OF POLYOMAVIRUS LATE PROMOTER ACTIVITY; ACTIVATION OF GENE-EXPRESSION) -more-? Display 6/9/3 (Item 1 from file: 434) DIALOG(R) File 434: SciSearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. (MURINE HEMATOPOIETIC PROGENITOR CELLS FOLLOWING RETROVIRAL 86-0127 001 TRANSFER; RETROVIRUS VECTORS; EXPRESSION OF HUMAN ADENOSINE-DEAMINASE) (RAT AORTIC SMOOTH-MUSCLE CELLS; HEAT-SHOCK PROTEINS; INVIVO BIOSYNTHESIS; EARLY MOUSE EMBRYO; TWO-DIMENSIONAL ELECTROPHORESIS; TYPE-III PROCOLLAGEN)

86-4753 001 (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG REGION)

86-5967 001 (SIMIAN VIRUS-40 REPLICATION; EXPRESSION IN COS CELLS; LATE GENE; PLASMID DNA; HOMOLOGOUS RECOMBINATION IN MAMMALIAN-CELLS; SV40 MUTANTS)

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Display 6/9/4 (Item 2 from file: 434) DIALOG(R)File 434:SciSearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv.

08028445 Genuine Article#: G9164 Number of References: 41
Title: EXPRESSION OF HUMAN CLASS-II MAJOR HISTOCOMPATIBILITY COMPLEX
ANTIGENS USING RETROVIRUS **VECTORS**

Author(s): KORMAN AJ; FRANTZ JD; STROMINGER JL; MULLIGAN RC Corporate Source: MIT, WHITEHEAD INST BIOMED RES/CAMBRIDGE//MA/02142; HARVARD UNIV, DEPT BIOCHEM & MOLEC BIOL/CAMBRIDGE//MA/02138

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Journal: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED
    STATES OF AMERICA, 1987, V84, N8, P2150-2154
                    Document Type: ARTICLE
Language: ENGLISH
Geographic Location: USA
Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences
Journal Subject Category: MULTIDISCIPLINARY SCIENCES
Research Fronts: 86-0127 003
                               (MURINE HEMATOPOIETIC PROGENITOR CELLS
    FOLLOWING RETROVIRAL TRANSFER; RETROVIRUS VECTORS; EXPRESSION OF
    HUMAN ADENOSINE-DEAMINASE)
               (HUMAN MAJOR HISTOCOMPATIBILITY COMPLEX; HLA CLASS-II
  86-1347 002
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      Display 6/9/4
                        (Item 2 from file: 434)
DIALOG(R) File 434: SciSearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
    ANTIGENS; DNA RESTRICTION FRAGMENT LENGTH POLYMORPHISMS;
    INSULIN-DEPENDENT DIABETES-MELLITUS)
                (TRANSCRIPTIONAL ENHANCER; VIRAL REGULATORY ELEMENTS;
  86-1674 001
    ADENOVIRUS E1A GENE; REGULATION OF POLYOMAVIRUS LATE PROMOTER
    ACTIVITY; ACTIVATION OF GENE-EXPRESSION)
                (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS
  86-4753 001
    OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG
    REGION)
                (PARVALBUMIN EXPRESSION IN MAMMALIAN SKELETAL-MUSCLE;
  86-6423 001
    DIFFERENTIATION ANTIGENS; IMMUNOLOGICAL DETECTION; RABBIT
    SKELETAL-MUSCLES; NEURAL CONTROL)
  86-7866 001
                (MOUSE MAMMARY-TUMOR CELLS; DNA TRANSFECTION; TRANSFECTED
    HUMAN-FIBROBLASTS; VIRAL VECTORS; ENHANCER FUNCTION; SINGLE
    PLASMID; GENETIC COMPLEMENTATION)
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86-4753 001 (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG REGION)

86-6423 001 (PARVALBUMIN EXPRESSION IN MAMMALIAN SKELETAL-MUSCLE; DIFFERENTIATION ANTIGENS; IMMUNOLOGICAL DETECTION; RABBIT SKELETAL-MUSCLES; NEURAL CONTROL)

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07847954 Genuine Article#: F6673 Number of References: 74
Title: EPSTEIN-BARR VIRUS GLYCOPROTEIN HOMOLOGOUS TO HERPES-SIMPLEX VIRUS-GB

Author(s): GONG M; OOKA T; MATSUO T; KIEFF E

Corporate Source: UNIV CHICAGO, KOVLER VIRAL ONCOL LABS/CHICAGO//IL/60637

Journal: JOURNAL OF VIROLOGY, 1987, V61, N2, P499-508

Language: ENGLISH Document Type: ARTICLE

Geographic Location: USA

Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences

Journal Subject Category: VIROLOGY

Research Fronts: 86-1382 002 (HEMAGGLUTININ NEURAMINIDASE GENE OF NEWCASTLE-DISEASE VIRUS; NUCLEOTIDE-SEQUENCE ANALYSIS; TRANSCRIPTION OF GENES; CDNA CLONE)

86-4807 002 (EPSTEIN-BARR VIRUS; ACQUIRED IMMUNE DEFICIENCY SYNDROME-RELATED LYMPHOPROLIFERATIVE DISORDERS; VIRUS-IMMORTALIZED LYMPHOBLASTOID CELL-LINES)

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- 86-0127 001 (MURINE HEMATOPOIETIC PROGENITOR CELLS FOLLOWING RETROVIRAL TRANSFER; RETROVIRUS **VECTORS**; EXPRESSION OF HUMAN ADENOSINE-DEAMINASE)
- 86-1455 001 (ESCHERICHIA-COLI K-12; MUTATIONS IN THE ARAC REGULATORY GENE; COOPERATIVE REGULATION OF UIDA EXPRESSION)
- 86-1491 001 (EPSTEIN-BARR VIRUS; CHRONIC INFECTIOUS MONONUCLEOSISLIKE SYNDROME IN COMMON MARMOSETS; PERSISTENT SYMPTOMS FOLLOWING INFECTIOUS-MONONUCLEOSIS)
- 86-1674 001 (TRANSCRIPTIONAL ENHANCER; VIRAL REGULATORY ELEMENTS; ADENOVIRUS E1A GENE; REGULATION OF POLYOMAVIRUS LATE PROMOTER ACTIVITY; ACTIVATION OF GENE-EXPRESSION)
- 86-4753 001 (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG REGION)
- 86-6423 001 (PARVALBUMIN EXPRESSION IN MAMMALIAN SKELETAL-MUSCLE; DIFFERENTIATION ANTIGENS; IMMUNOLOGICAL DETECTION; RABBIT SKELETAL-MUSCLES; NEURAL CONTROL)

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86-6701 001 (OLIGOSACCHARIDE PROCESSING; PROTEIN GLYCOSYLATION; SECRETORY PROTEINS; MUNG BEAN SEEDLINGS; RAT-LIVER ALPHA-MANNOSIDASE; N-LINKED CARBOHYDRATE)

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TYPE-IV COLLAGEN)

- 86-1382 001 (HEMAGGLUTININ NEURAMINIDASE GENE OF NEWCASTLE-DISEASE VIRUS; NUCLEOTIDE-SEQUENCE ANALYSIS; TRANSCRIPTION OF GENES; CDNA CLONE)
- 86-1674 001 (TRANSCRIPTIONAL ENHANCER; VIRAL REGULATORY ELEMENTS; ADENOVIRUS E1A GENE; REGULATION OF POLYOMAVIRUS LATE PROMOTER ACTIVITY; ACTIVATION OF GENE-EXPRESSION)
- 86-2124 001 (GLIAL FIBRILLARY ACIDIC PROTEIN; CENTRAL NERVOUS-SYSTEM TUMORS; GROWTH OF HUMAN OLIGODENDROCYTES INVITRO; ASTROCYTES IN PRIMARY CULTURES)
- 86-4753 001 (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG REGION)
- 86-6423 001 (PARVALBUMIN EXPRESSION IN MAMMALIAN SKELETAL-MUSCLE; DIFFERENTIATION ANTIGENS; IMMUNOLOGICAL DETECTION; RABBIT

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SKELETAL-MUSCLES; NEURAL CONTROL)

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                                     Number of References: 62
07602552
           Genuine Article#: E5673
Title: EXPRESSION OF A MOLECULARLY CLONED HUMAN C-SRC ONCOGENE BY USING A
    REPLICATION-COMPETENT RETROVIRAL VECTOR
Author(s): TANAKA A; FUJITA DJ
Corporate Source: UNIV WESTERN ONTARIO, CANC RES LAB/LONDON N6A
    5B7/ONTARIO/CANADA/; UNIV WESTERN ONTARIO, DEPT BIOCHEM/LONDON N6A
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Journal: MOLECULAR AND CELLULAR BIOLOGY, 1986, V6, N11, P3900-3909
Language: ENGLISH
                    Document Type: ARTICLE
Geographic Location: CANADA
Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences
Journal Subject Category: BIOCHEMISTRY & MOLECULAR BIOLOGY
Research Fronts: 86-3403 003
                               (ROUS-SARCOMA VIRUS; TYROSINE
    PHOSPHORYLATION; TRANSFORMING PROTEIN; PP60C-SRC KINASE; EPIDERMAL
    GROWTH-FACTOR RECEPTOR)
               (RAS PROTEINS; CONTINUED EXPRESSION OF THE C-HA-RAS
  86-3728 002
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   ONCOGENE; TRANSFORMING GENES IN HUMAN-LEUKEMIA CELLS)
  86-1382 001
                (HEMAGGLUTININ NEURAMINIDASE GENE OF NEWCASTLE-DISEASE
    VIRUS; NUCLEOTIDE-SEQUENCE ANALYSIS; TRANSCRIPTION OF GENES; CDNA
   CLONE)
  86-1674 001
                (TRANSCRIPTIONAL ENHANCER; VIRAL REGULATORY ELEMENTS;
   ADENOVIRUS E1A GENE; REGULATION OF POLYOMAVIRUS LATE PROMOTER
   ACTIVITY; ACTIVATION OF GENE-EXPRESSION)
  86-4303 001
                (PLATELET-DERIVED GROWTH-FACTOR; RECEPTOR PROTO-ONCOGENE
    FMS; TRANSFORMING GROWTH-FACTORS)
  86-4753 001
                (ROUS-SARCOMA VIRUS SRC GENE; HOST RANGE DETERMINANTS
   OF AVIAN RETROVIRUS ENVELOPE GENES; AMINO-ACID SEQUENCE HOMOLOGY IN GAG
   REGION)
                (PARVALBUMIN EXPRESSION IN MAMMALIAN SKELETAL-MUSCLE;
  86-6423 001
    DIFFERENTIATION ANTIGENS; IMMUNOLOGICAL DETECTION; RABBIT
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